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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/677,807	10/01/2003	David Epstein	23239-540 (ARC-40)	9987	
30623	7590 03/14/2006		EXAMINER		
•	IN, COHN, FERRIS, G	WOLLENBERGER, LOUIS V			
AND POPEO, P.C. ONE FINANCIAL CENTER			ART UNIT	PAPER NUMBER	
BOSTON, MA	BOSTON, MA 02111			1635	
			DATE MAILED: 03/14/2006	5 .	

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)	
	10/677,807	EPSTEIN ET AL.	
Office Action Summary	Examiner	Art Unit	
	Louis V. Wollenberger	1635	
The MAILING DATE of this communication	appears on the cover sheet with t	he correspondence address	
Period for Reply  A SHORTENED STATUTORY PERIOD FOR REWHICHEVER IS LONGER, FROM THE MAILING  - Extensions of time may be available under the provisions of 37 CF after SIX (6) MONTHS from the mailing date of this communication. If NO period for reply is specified above, the maximum statutory period for reply within the set or extended period for reply will, by some Any reply received by the Office later than three months after the meanned patent term adjustment. See 37 CFR 1.704(b).	G DATE OF THIS COMMUNICAT R 1.136(a). In no event, however, may a reply n. eriod will apply and will expire SIX (6) MONTHS tatute, cause the application to become ABAND	FION.  be timely filed  from the mailing date of this communication.  DONED (35 U.S.C. § 133).	
Status			
Responsive to communication(s) filed on <i>Q</i> This action is <b>FINAL</b> . 2b) ☑      Since this application is in condition for all closed in accordance with the practice und	This action is non-final.  Dwance except for formal matters		
Disposition of Claims			
4)  Claim(s) 1-7 is/are pending in the applicating 4a) Of the above claim(s) is/are with 5)  Claim(s) is/are allowed.  6)  Claim(s) is/are rejected.  7)  Claim(s) is/are objected to.  8)  Claim(s) 1-7 are subject to restriction and/or	drawn from consideration.		
Application Papers			
9) The specification is objected to by the Exart 10) The drawing(s) filed on is/are: a) Applicant may not request that any objection to Replacement drawing sheet(s) including the count of the county The oath or declaration is objected to by the second se	accepted or b) objected to by the drawing(s) be held in abeyance.  prection is required if the drawing(s) in	See 37 CFR 1.85(a). s objected to. See 37 CFR 1.121(d).	
Priority under 35 U.S.C. § 119			
<ul> <li>12) Acknowledgment is made of a claim for forma) All b) Some * c) None of:</li> <li>1. Certified copies of the priority documents.</li> <li>2. Certified copies of the priority documents.</li> <li>3. Copies of the certified copies of the application from the International But * See the attached detailed Office action for a second content.</li> </ul>	nents have been received. nents have been received in Appl priority documents have been rec ireau (PCT Rule 17.2(a)).	ication No ceived in this National Stage	
Attachment(s)  1) Notice of References Cited (PTO-892)  2) Notice of Draftsperson's Patent Drawing Review (PTO-948  3) Information Disclosure Statement(s) (PTO-1449 or PTO/SI Paper No(s)/Mail Date	Paper No(s)/M	mary (PTO-413) ail Date mal Patent Application (PTO-152)	

# DETAILED ACTION

## Election/Restrictions

Restriction to one of the following inventions is required under 35 U.S.C. 121:

- I. Claims 1 and 2, drawn to an aptamer and to a therapeutic composition for treating subjects having HIV infection, classified in class 536, subclass 24.5. Election of this group requires the further election of a single aptamer sequence from claim 1 and 2, as explained below.
- II. Claims 3, 4, 6, and 7, drawn to a method of treating HIV infection in a subject using an aptamer, classified in class 514, subclass 44, for example. Election of this group requires the further election of a single aptamer sequence from claim 3, as explained below.
- III. Claim 5, drawn to a method for diagnosing HIV infection, comprising the use of an aptamer-reporter complex, classified in class 435, subclass 6. Election of this group requires the further election of a single aptamer sequence from claim 5, as explained below.

The inventions are distinct, each from the other because of the following reasons:

Invention I is related to inventions II and III as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product. See MPEP § 806.05(h). In the instant case the processes as claimed can be practiced

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with another materially different product. For example, the method of treating HIV infected subjects can be accomplished with small molecule, non-nucleic acid reagents or with antisense or siRNA compounds, which does not require the use of an aptamer such as that claimed in Group I. The method for diagnosing HIV infection can be accomplished with a suitable antibody or by using quantitative PCR, which, similarly, does not require the use of an aptamer.

Inventions II and III are unrelated. Inventions are unrelated if it can be shown that they are not disclosed as capable of use together and they have different designs, modes of operation, and effects (MPEP § 802.01 and § 806.06). In the instant case the different inventions are unrelated because they are drawn to methods that are distinct both physically and functionally, and are not required one for the other. The different methods require different steps and/or use different molecules. For example, Group III requires the use of an aptamer-reporter complex and requires steps for providing a sample from a subject, contacting the sample with the aptamerreporter complex, and would also require measuring and/or quantifying the signal relative to a suitable reference, which steps and reagents are not required by Group II. Group II requires a step for administering the aptamer to a subject, which is not a step required by Group III.

In addition, each group would require different keyword searches (i.e., different fields of search) and different considerations of the patent and non-patent literature with regard to novelty and unobviousness.

Therefore, because these inventions are distinct for the reasons given above, and the searches required for each are divergent and not coextensive, and because a search and examination of Inventions I, II, and III in a single application would present a serious burden on the Examiner, restriction for examination purposes as indicated is proper.

## Restriction to a single aptamer sequence SEQ ID NO:

Should applicants elect to prosecute Group I, II, or III, each of these groups is subject to further restriction as follows.

Claims 1, 2, 3, and 5 recite a plurality of different aptamer sequences, which may be used for treating and/or diagnosing HIV infection. Specifically, the instant claims recite Markush groups consisting of several different SEQ ID Nos.

Pursuant to 35 U.S.C. 121 and 37 C.F.R. 1.141, these sequences (i.e., SEQ ID Nos) are subject to restriction since they are not considered to be a proper genus/Markush. See MPEP 803.02 - PRACTICE RE MARKUSH-TYPE CLAIMS - If the members of the Markush group are sufficiently few in number or so closely related that a search and examination of the entire claim can be made without serious burden, the examiner must examine all the members of the Markush group in the claim on the merits, even though they are directed to independent and distinct inventions. In such a case, the examiner will not follow the procedure described below and will not require restriction. Since the decisions in *In re Weber*, 580 F.2d 455, 198 USPQ 328 (CCPA 1978) and *In re Haas*, 580 F.2d 461, 198 USPQ 334 (CCPA 1978), it is improper for the Office to refuse to examine that which applicants regard as their invention, unless the subject matter in a claim lacks unity of invention. *In re Harnish*, 631 F.2d 716, 206 USPQ 300 (CCPA 1980); and *Ex parte Hozumi*, 3 USPQ2d 1059 (Bd. Pat. App. & Int. 1984).

Broadly, unity of invention exists where compounds included within a Markush group (1) share a common utility, and (2) share a substantial structural feature disclosed as being essential to that utility.

In the instant case, the sequences do not appear to share a substantial structural feature disclosed as being essential to a common utility. That is, the recited sequences appear to be structurally and functionally distinct, one from the other, and are therefore considered to be independent and distinct inventions. Although the different sequences may target and bind the same polypeptide, the different sequences may differ with regard to binding specificity and affinity, for example, which may confer distinct properties to the claimed invention as a whole.

Furthermore, a search and examination of more than one (1) of the sequences claimed in the instant claims presents an undue burden on the Patent and Trademark Office due to the complex nature of the search in terms of computer time needed to perform the search and the subsequent analysis of the search results by the examiner. In view of the foregoing, one (1) aptamer sequence is considered to be a reasonable number of sequences for examination.

Accordingly, applicants are required to elect one (1) sequence SEQ ID NO: from the instant claims for prosecution on the merits with the elected group. Note that this is not a species election.

In the event of rejoinder of product and process claims, applicants are reminded that the process claims identified above depending from the product claims identified above must recite the same SEQ ID No to remain consonant with this restriction requirement.

## Rejoinder

The examiner has required restriction between product and process claims. Where applicant elects claims directed to the product, and a product claim is subsequently found allowable, withdrawn process claims that depend from or otherwise include all the limitations of the allowable product claim will be rejoined in accordance with the provisions of MPEP §

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821.04. Process claims that depend from or otherwise include all the limitations of the patentable product will be entered as a matter of right if the amendment is presented prior to final rejection or allowance, whichever is earlier. Amendments submitted after final rejection are governed by 37 CFR 1.116; amendments submitted after allowance are governed by 37 CFR 1.312.

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In the event of rejoinder, the requirement for restriction between the product claims and the rejoined process claims will be withdrawn, and the rejoined process claims will be fully examined for patentability in accordance with 37 CFR 1.104. Thus, to be allowable, the rejoined claims must meet all criteria for patentability including the requirements of 35 U.S.C. 101, 102, 103, and 112. Until an elected product claim is found allowable, an otherwise proper restriction requirement between product claims and process claims may be maintained. Withdrawn process claims that are not commensurate in scope with an allowed product claim will not be rejoined. See "Guidance on Treatment of Product and Process Claims in light of In re Ochiai, In re Brouwer and 35 U.S.C. § 103(b)," 1184 O.G. 86 (March 26, 1996). Additionally, in order to retain the right to rejoinder in accordance with the above policy, Applicant is advised that the process claims should be amended during prosecution either to maintain dependency on the product claims or to otherwise include the limitations of the product claims. Failure to do so may result in a loss of the right to rejoinder. Further, note that the prohibition against double patenting rejections of 35 U.S.C. 121 does not apply where the restriction requirement is withdrawn by the examiner before the patent issues. See MPEP § 804.01.

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#### Conclusion

Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Louis V. Wollenberger whose telephone number is 571-272-8144. The examiner can normally be reached on Mon–Fri, 8:00 am–4:30 pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's acting supervisor, Andrew Wang can be reached on 571-272-0811. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval system (PAIR). Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

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For all other customer support, please call the USPTO Call Center (UCC) at 800-786-9199.

SEAN McGARRY PRIMARY EXAMINER Louis V. Wollenberger, Ph.D.

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Examiner

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March 6, 2006